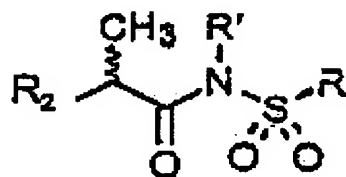


CLAIMS

1. Use of N-(2-aryl-propionyl)-sulfonamides of general formula (I):



(I)

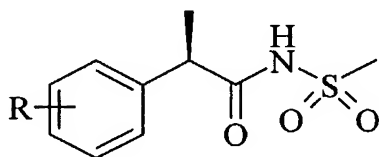
5 in which

R₂ is an aryl group,

R is a straight or branched C₁-C₆-alkyl, trifluoromethyl, cyclohexyl, o-tolyl, 3-pyridyl, 2-pyridyl-ethyl, p-cyano-phenylmethyl, p-aminophenylmethyl, 3-cyano-1-propyl, 4-aminobutyl group, an alkoxyethylene CH₃-(CH₂)_{n_i}-
 10 (OCH₂CH₂)_{m_i}- group in which n_i is zero or 1 and m_i is an integer 1 to 3, or a P₁P₂N-CH₂-CH₂- group in which P₁ and P₂ are independently H, C₁-C₃- alkyl, benzyloxy-carbonyl, α-, β- or α-pyridocarbonyl, carboxycarbonyl or carbalkoxycarbonyl, or P₁ and P₂, when joined to the N atom which they are linked to, form a phthalimido, piperidino, morpholino residue;

15 R' is H or straight or branched C₁-C₃-alkyl, preferably hydrogen, for the preparation of a medicament for the treatment of spinal cord injury.

2. Use according to claim 1 of the compounds of formula (Ia)



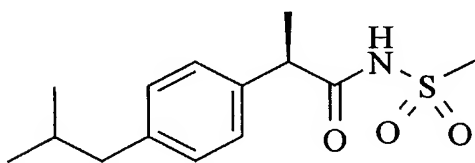
(Ia)

20

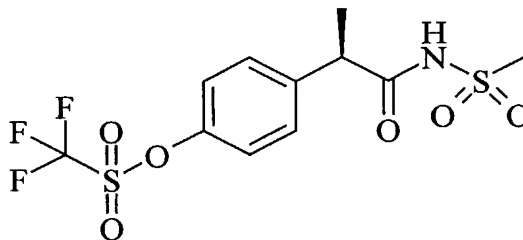
wherein R represents one to three substituents, which are the same or different, selected from hydrogen, halogen atoms, C₁-C₄-alkyl, C₁-C₄-alkoxy,

hydroxy, C₁-C₇-acyloxy, cyano, nitro, amino, C₁-C₃-acylamino, halo C₁-C₃-alkyl, halo C₁-C₃-alkoxy, benzoyl, 4-(2-methyl-propyl)-phenyl, 3-phenoxy-phenyl, 2-[4-(1-oxo-2-isoindoliny)]phenyl, 5-benzoyl-thien-2-yl, 4-thienoyl-phenyl, C₁-C₂-halogenoalkylsulphonyloxy.

- 5 3. Use according to claim 2 wherein R represents hydrogen, 4-isobutyl, 3-benzoyl, 4-trifluoromethanesulphonyloxy.
4. Use according to claim 2 of the compounds of formula (II) and (III).



(II)



(III)